

**IN THE UNITED STATES PATENT AND TRADEMARK OFFICE**

In re Application of

**Paul Habermann**

Application No.: **10/076,631**

Filed: **February 19, 2002**

Title: **NUCLEIC ACIDS ENCODING A  
HIRUDIN AND PRO-INSULIN AS  
SUPERSCRETABLE PEPTIDES  
AND FOR PARALLEL  
IMPROVEMENT OF THE  
EXPORTED FORMS OF ONE OR  
MORE POLYPEPTIDES OF  
INTEREST**

Examiner: **KOSSON,  
Roseanne**

Art Unit: **1652**

Confirmation **2601**  
No.

**CERTIFICATE OF EFS-WEB TRANSMISSION**

I hereby certify that the correspondence below is being transmitted via the USPTO's electronic filing system in accordance with 1.6(a)(4), on the date indicated below.

Date of Deposit July 9, 2010

Printed Name of Person

Signing Certificate Delia Coughlin

Signature /Delia Coughlin/

**REQUEST FOR CERTIFICATE OF CORRECTION**  
**UNDER 35 U.S.C. 254/255 and 37 C.F.R. 1.322/323**

Commissioner for Patents  
Attn: Certificate of Correction Branch  
P. O. Box 1450  
Alexandria, VA 22313-1450

The following is a request for a certificate of correction in Serial Number 10/076,631, now patent Number 7,638,618.

A certificate of correction under 35 U.S.C. 254 is respectfully requested in the above-identified patent.

- ☐ All errors were the fault of the USPTO, no fee required. In the event that a further fee is required, please charge the amount to Deposit Account No. **18-1982**.
- ☐ All errors were the fault of the applicant and, accordingly, please charge **\$100.00** to our Deposit Account No. **18-1982**. In the event that a further fee is required, please charge the amount to the same Deposit Account.
- ☒ The errors were the fault of both the applicant and the USPTO and, accordingly, please charge **\$100.00** to our Deposit Account No. **18-1982**. In the event that a further fee is required, please charge the amount to the same Deposit Account.

The exact location where the error appears in the patent and patent application is noted below.

The requested correction is attached on Form PTO 1050.

**EXACT LOCATION WHERE ERRORS APPEAR**

1. In column 7, line 55 (Page 3, Amendments to the Specification, (05/19/2006), line 31): “(Refludan®)” should read as - - (REFLUDAN®) - -.
2. In column 8, line 2 (Page 17, Specification, (02/19/2002), line 16): “hirF1” should read as - - hIRF1 - -.
3. In column 9, line 67 (Page 4, Amendments to the Specification, (05/19/2006), line 4): “REFLUDAN®.” should read as - - REFLUDAN® - -.
4. In column 10, line 10 (Page 4, Amendments to the Specification, (05/19/2006), line 7): “INVITROGEN®.” should read as - - INVITROGEN® - -.
5. In column 10, line 30 (Page 23, Specification, (02/19/2002), line 9): “primer Pichia” should read as - - Primer pichia - -.
6. In column 10, line 46 (Page 23, Specification, (02/19/2002), line 20): “zeocine” should read as - - zeocin - -.
7. In Claim 6, (see Claim 10 as presented by Amendment filed Jul. 16, 2009): column 16, line 50: “factis,” should read as - - lactis, - -.

Respectfully submitted,

/George S Jones/

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Sanofi-aventis U.S. Docket No. DEAV2001/0007  
Date: July 8, 2010

## UNITED STATES PATENT AND TRADEMARK OFFICE CERTIFICATE OF CORRECTION

PATENT NO. : 7,638,618

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APPLICATION NO.: 10/076,631

ISSUE DATE : December 29, 2009

INVENTOR(S) : Paul Habermann

It is certified that an error appears or errors appear in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

In column 7, line 55, delete "(Refludan®)" and insert - - (REFLUDAN®) - -, therefor.

In column 8, line 2, delete "hirF1" and insert - - hIRF1 - -, therefor.

In column 9, line 67, delete "REFLUDAN®." and insert - - REFLUDAN® - -, therefor.

In column 10, line 10, delete "INVITROGEN®." and insert - - INVITROGEN® - -, therefor.

In column 10, line 30, delete "primer Pichia" and insert - - Primer pichia - -, therefor.

In column 10, line 46, delete "zeocine" and insert - - zeocin - -, therefor.

In column 16, line 50, in Claim 6, delete "factis," and insert - - lactis, - -, therefor.

MAILING ADDRESS OF SENDER (Please do not use customer number below)

This collection of information is required by 37 CFR 1.322, 1.323, and 1.324. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.14. This collection is estimated to take 1.0 hour to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, P.O. Box 1450, Alexandria, VA 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. **SEND TO: Attention Certificate of Corrections Branch, Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.**

*If you need assistance in completing the form, call 1-800-PTO-9199 and select option 2*

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The information provided by you in this form will be subject to the following routine uses:

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6. A record in this system of records may be disclosed, as a routine use, to another federal agency for purposes of National Security review (35 U.S.C. 181) and for review pursuant to the Atomic Energy Act (42 U.S.C. 218(c)).
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8. A record from this system of records may be disclosed, as a routine use, to the public after either publication of the application pursuant to 35 U.S.C. 122(b) or issuance of a patent pursuant to 35 U.S.C. 151. Further, a record may be disclosed, subject to the limitations of 37 CFR 1.14, as a routine use, to the public if the record was filed in an application which became abandoned or in which the proceedings were terminated and which application is referenced by either a published application, an application open to public inspection or an issued patent.
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